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GenCore version 5.1.3
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OM protein - protein search, using sw model

Run on: November 9, 2002, 03:16:01 : Search time 80 Seconds
(without alignments)
316.470 Million cell updates/sec

Title: US-09-895-298A-83
Perfect score: 1002
Sequence: 1 MMNFQPPSKAMRASQMTFF.....HDSLIDLRSSRVQEGNPRA 190

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq.101002.*

- 1: /SIDS2/gcgdata/geneseq-emb1/AA1980.DAT:*
- 2: /SIDS2/gcgdata/geneseq-emb1/AA1981.DAT:*
- 3: /SIDS2/gcgdata/geneseq-emb1/AA1982.DAT:*
- 4: /SIDS2/gcgdata/geneseq-emb1/AA1983.DAT:*
- 5: /SIDS2/gcgdata/geneseq-emb1/AA1984.DAT:*
- 6: /SIDS2/gcgdata/geneseq-emb1/AA1985.DAT:*
- 7: /SIDS2/gcgdata/geneseq-emb1/AA1986.DAT:*
- 8: /SIDS2/gcgdata/geneseq-emb1/AA1987.DAT:*
- 9: /SIDS2/gcgdata/geneseq-emb1/AA1988.DAT:*
- 10: /SIDS2/gcgdata/geneseq-emb1/AA1989.DAT:*
- 11: /SIDS2/gcgdata/geneseq-emb1/AA1990.DAT:*
- 12: /SIDS2/gcgdata/geneseq-emb1/AA1991.DAT:*
- 13: /SIDS2/gcgdata/geneseq-emb1/AA1992.DAT:*
- 14: /SIDS2/gcgdata/geneseq-emb1/AA1993.DAT:*
- 15: /SIDS2/gcgdata/geneseq-emb1/AA1994.DAT:*
- 16: /SIDS2/gcgdata/geneseq-emb1/AA1995.DAT:*
- 17: /SIDS2/gcgdata/geneseq-emb1/AA1996.DAT:*
- 18: /SIDS2/gcgdata/geneseq-emb1/AA1997.DAT:*
- 19: /SIDS2/gcgdata/geneseq-emb1/AA1998.DAT:*
- 20: /SIDS2/gcgdata/geneseq-emb1/AA1999.DAT:*
- 21: /SIDS2/gcgdata/geneseq-emb1/AA2000.DAT:*
- 22: /SIDS2/gcgdata/geneseq-emb1/AA2001.DAT:*
- 23: /SIDS2/gcgdata/geneseq-emb1/AA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	1002	100.0	191	21	AAB24458	Human secreted pro
2	1002	100.0	191	22	AAB83082	Human CASB6411-rel
3	1002	100.0	268	22	AAM79104	Human protein SEQ
4	1002	100.0	280	22	ABB11361	Human LAK-4p homol
5	1002	100.0	280	22	AAM80088	Human protein SEQ
6	1002	100.0	330	22	AAB95481	Human protein sequ
7	1002	100.0	387	21	AAB08764	A human leukocyte
8	1002	100.0	438	22	AAB83081	Human CASB6411-rel
9	1002	100.0	460	22	AAB83079	Human CASB6411 pro
10	292.5	29.2	305	23	ABP41828	Human ovarian anti

11	148	14.8	31	22	ABB39891	Peptide #7397 enco
12	148	14.8	31	22	AAM60631	Human brain expres
13	148	14.8	31	22	AAM73303	Human bone marrow
14	148	14.8	31	22	AAM33503	Peptide #7540 enco
15	148	14.8	31	23	ABG43154	Human peptide enco
16	122	12.2	88	20	AA100264	Human secreted pro
17	97	9.7	992	22	ABG28109	Novel human diagno
18	94	9.4	40	20	AA100348	Fragment of human
19	90.5	9.0	229	23	ABG64734	Human albumin fusi
20	90.5	9.0	229	23	ABG64736	Human albumin fusi
21	90.5	9.0	229	23	AAO17174	Human secreted pro
22	90.5	9.0	229	23	AAO17199	Human secreted pro
23	90.5	9.0	334	23	AAO17232	Human secreted pro
24	89	8.9	272	21	AAO14398	Arabidopsis thalia
25	89	8.9	291	21	AAO14397	Arabidopsis thalia
26	88.5	8.8	589	21	AAO14396	Human secreted pro
27	88.5	8.8	706	22	AAO14395	Human membrane ass
28	87	8.7	272	21	AAO14394	Arabidopsis thalia
29	87	8.7	291	21	AAO14393	Arabidopsis thalia
30	84	8.4	273	21	AAO14392	Arabidopsis thalia
31	83.5	8.3	339	23	ABP26928	Streptococcus poly
32	83.5	8.3	339	23	ABP29763	Streptococcus poly
33	82.5	8.2	308	22	AAO14391	S. epidermidis ope
34	82.5	8.2	310	22	ABP28118	Mouse T2R31 amino
35	80.5	8.0	248	23	ABP28117	Human polypeptide
36	80	8.0	203	23	ABP28116	Streptococcus poly
37	80	8.0	242	21	AAO14390	Pinus radiata ane
38	80	8.0	426	22	AAO14389	Human PRO6028 poly
39	79.5	7.9	154	23	ABP28115	Listeria monocytog
40	79	7.9	473	22	AAO14388	S. epidermidis ope
41	79	7.9	496	23	ABP38156	Staphylococcus epi
42	78	7.8	107	18	AAO14387	S. pneumoniae prot
43	77.5	7.7	306	21	AAO14386	Plasmodium falci
44	77.5	7.7	596	22	AAO14385	Staphylococcus aur
45	77.5	7.7	604	22	AAO14384	Staphylococcus aur

ALIGNMENTS

RESULT 1	
AAB24458	
ID AAB24458 standard; Protein; 191 AA.	
XX	
AC AAB24458;	
XX	
DT 20-NOV-2000 (first entry)	
XX	
DE Human secreted protein sequence encoded by gene 22 SEQ ID NO:83.	
XX	
KW Human; secreted protein; cytosolic; antianaemic; antidiabetic;	
KW antinflammatory; ophthalmological; antirheumatic; antiarthritic;	
KW antipsoriatic; antiangiogenic; cardiant; anti-HIV; nootropic;	
KW neuroprotective; antimicrobial; antiparkinsonian; cancer;	
KW immune system disorder; angiogenesis; hyperproliferative disorder;	
KW cardiovascular disorder; apoptosis; neurological disease;	
KW infectious disease; wound healing.	
XX	
OS Homo sapiens.	
XX	
PN WO200035937-A1.	
XX	
PD 22-JUN-2000.	
XX	
PF 16-DEC-1999; 99WO-US29950.	
XX	
PR 17-DEC-1998; 98US-0112809.	
XX	
PA 18-DEC-1998; 98US-0113006.	
XX	
PI (HUMA-) HUMAN GENOME SCI INC.	
PI Ruben SM, Ebner R, Rosen CA, Endress CA, Soppet DR, Ni J;	
PI Duan DR, Moore PA, Shi Y, Lafleur DW, Olsen HS, Florence K;	

```
XX WPI; 2000-431566/37.
DR N-PSDB; AAA78402.
XX
PT Forty seven human nucleic acids encoding secreted proteins, useful in
PT the treatment, prevention and diagnosis of cancers, disorders of the
PT immune system, angiogenesis disorders, neurological diseases and
PT hyperproliferative disorders
XX
PS Claim 11; Page 496; 562pp; English.
XX
CC The polynucleotide sequence given in AAA78381 to AAA78432 encode the
CC human secreted proteins given in AAB24437 to AAB24604. Human secreted
CC proteins have activities based on the tissues and cells the genes are
CC expressed in. Examples of activities include: cytostatic; antianaemic;
CC antidiabetic; antiinflammatory; ophthalmological; antitumumatic;
CC antiarthritic; antipsoriatic; antiangiogenic; cardiant; anti-HIV;
CC neurotropic; neuroprotective; antimicrobial and antiparkinsonian.
CC Human secreted protein polynucleotides, polypeptides, antagonists and/or
CC agonists may be useful in treating, preventing, and/or diagnosing other
CC diseases, disorders, and/or conditions such as: (a) cancers; (b)
CC disorders of the immune system; (c) angiogenesis disorders; (d)
CC hyperproliferative disorders; (e) cardiovascular disorders; (f) diseases
CC associated with increase apoptosis; (g) neurological diseases; and
CC (h) infectious diseases. They are also used to promote wound healing.
CC AAA78372 to AAA78380 and AAB24436 represent sequences used in the
CC exemplification of the present invention.
XX
SQ Sequence 191 AA;

Query Match 100.0%; Score 1002; DB 21; Length 191;
Best Local Similarity 100.0%; Pred. No. 2.2e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MMNFQPPSKAWRASQMTFFIFLFFPSFTGYLCTLATITWRLKPSADCGPFGPLPIFIH 60
DB 1 MMNFQPPSKAWRASQMTFFIFLFFPSFTGYLCTLATITWRLKPSADCGPFGPLPIFIH 60
QY 61 SIYSWIDTLSTRPGYLWVWVIYRNLIIGSVHFFILTLVLIITYLYWQITGGRKIMIRLL 120
DB 61 SIYSWIDTLSTRPGYLWVWVIYRNLIIGSVHFFILTLVLIITYLYWQITGGRKIMIRLL 120
QY 121 HEQIINEGKDKMFLEIKLIKQDMKKANPSSLVLERREVEQOGFLHGEHDSLDLSR 180
DB 121 HEQIINEGKDKMFLEIKLIKQDMKKANPSSLVLERREVEQOGFLHGEHDSLDLSR 180
QY 181 RSVQEGNPRA 190
DB 181 RSVQEGNPRA 190

RESULT 2
AAB83082
ID AAB83082 standard; Protein; 191 AA.
XX
AC AAB83082;
XX
DT 29-JUN-2001 (first entry)
XX
DE Human CASB6411-related partial polypeptide #2.
XX
KW Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;
KW ovarian cancer; colon cancer; autoimmune disease.
XX
OS Homo sapiens.
XX
PN WO200123417-A2.
XX
PD 05-APR-2001.
XX
PF 27-SEP-2000; 2000MO-EP095500.
XX
PR 30-SEP-1999; 99GB-0023154.
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PR 07-JUL-2000; 2000GB-0016839.
XX
PA (SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.
XX
PI Vinals De Bassols YC;
XX
DR WPI; 2001-316133/33.
DR N-PSDB; AAF82463.
XX
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for
PT prophylactic and therapeutic treatment of cancers, particularly ovarian
PT and colon cancers, autoimmune diseases and related conditions
XX
PS Disclosure; Page 67; 95pp; English.
XX
CC The present sequence is provided in a specification relating
CC to CASB6411 polypeptides comprising a sequence having at least 70%
CC identity to a sequence of 460 or 154 amino acids fully defined in
CC the specification. CASB6411 polypeptides and polynucleotides are
CC useful for treating a subject by immunoprophylaxis or therapy.
CC The CASB6411 polypeptides are useful in diagnostics, and as
CC vaccines for prophylactic and therapeutic treatment of cancers,
CC particularly ovarian and colon cancers, autoimmune diseases and related
CC conditions. CASB6411 polypeptides are also useful for the
CC structure-based design of agonists, antagonists or inhibitors of the
CC polypeptide.
XX
SQ Sequence 191 AA;

Query Match 100.0%; Score 1002; DB 22; Length 191;
Best Local Similarity 100.0%; Pred. No. 2.2e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MMNFQPPSKAWRASQMTFFIFLFFPSFTGYLCTLATITWRLKPSADCGPFGPLPIFIH 60
DB 2 MMNFQPPSKAWRASQMTFFIFLFFPSFTGYLCTLATITWRLKPSADCGPFGPLPIFIH 61
QY 61 SIYSWIDTLSTRPGYLWVWVIYRNLIIGSVHFFILTLVLIITYLYWQITGGRKIMIRLL 120
DB 62 SIYSWIDTLSTRPGYLWVWVIYRNLIIGSVHFFILTLVLIITYLYWQITGGRKIMIRLL 121
QY 121 HEQIINEGKDKMFLEIKLIKQDMKKANPSSLVLERREVEQOGFLHGEHDSLDLSR 180
DB 122 HEQIINEGKDKMFLEIKLIKQDMKKANPSSLVLERREVEQOGFLHGEHDSLDLSR 181
QY 181 RSVQEGNPRA 190
DB 182 RSVQEGNPRA 191

RESULT 3
AAM79104
ID AAM79104 standard; Protein; 268 AA.
XX
AC AAM79104;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human protein SEQ ID NO 1766.
XX
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation.
XX
OS Homo sapiens.
XX
PN WO200157190-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001MO-US04098.
XX
```

PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.
PR 19-JUL-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;
XX
XX WPI: 2001-476283/51.
DR N-PSDB; AAK52237.
XX
XX Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -
PS Claim 20; Page 4113-4114; 6221pp; English.
XX
XX The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
CC inflammation.
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666
CC (AAM80020) are omitted as the relevant pages from the sequence listing
CC were missing at the time of publication.
XX
XX Sequence 268 AA:
SQ
Query Match 100.0%; Score 1002; DB 22; Length 268;
Best Local Similarity 100.0%; Pred. No. 3.4e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MMNFPPSKAMRARSQMMTFIFLFFPSTGVLCTLAITTWRLKPSADCGPFRGIPLFTH 60
DB 79 MMNFPPSKAMRARSQMMTFIFLFFPSTGVLCTLAITTWRLKPSADCGPFRGIPLFTH 138
QY 61 SIYSWIDTLSRRPGYLMWVWYIRNLIGSVHFFFLTLIVLITLYLWQTEGKIMIRLL 120
DB 139 SIYSWIDTLSRRPGYLMWVWYIRNLIGSVHFFFLTLIVLITLYLWQTEGKIMIRLL 198
QY 121 HEQIINEGKDKMFLIEKLKIQDMKKANPSSLVLERREVEOQGLHGEHDSIDLRSR 180
DB 199 HEQIINEGKDKMFLIEKLKIQDMKKANPSSLVLERREVEOQGLHGEHDSIDLRSR 258
QY 181 RSVQEGNPRA 190
DB 259 RSVQEGNPRA 268
RESULT 4
ABBI1361
ID ABBI1361 standard; peptide; 280 AA.
XX
XX ABB11361;
AC
XX 11-JAN-2002 (first entry)
DT
XX Human LAK-4p homologue, SEQ ID NO:1731.
DE
XX Human; cytokine; cell proliferation; cell differentiation; growth factor;
KW haematopoiesis regulation; tissue growth; immunomodulator; activin;

KW inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;
KW proliferation; metastasis; cancer; tumour; haematopoietic disorder;
KW myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;
KW chronic inflammatory condition; proliferative retinopathy;
KW atherosclerosis; coronary heart disease; arterial ischaemia;
KW bone disorder; osteoporosis; vascular growth disorder;
KW tissue regeneration; wound healing; infection; immune disorder;
KW cell culture; drug screening; gene therapy; antiinflammatory;
KW antiasthmatic; antiarthritic; haemostatic; antiarteriosclerotic;
KW cytosolic; osteopathic; vasotropic; cardiant; virucide; antibacterial;
KW antifungal; vulnerrary; antitumor.
XX
XX Homo sapiens.
XX
XX WO200157188-A2.
XX
XX 09-AUG-2001.
XX
XX 05-FEB-2001; 2001WO-US03800.
XX
XX 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Drmanac RT;
PI
XX
XX WPI: 2001-457740/49.
DR N-PSDB; ABA08605.
XX
XX Claim 20; Page 173; 1963pp; English.
XX
XX Sequences ABB10981-ABBI2330 represent 1350 novel human polypeptides, and
CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The
CC invention also relates to vectors and recombinant host cells comprising a
CC nucleotide of the invention, methods of producing the novel polypeptides,
CC antibodies against the polypeptides, methods of detecting the nucleotides
CC or polypeptides in a sample, and methods of identifying compounds which
CC bind to polypeptides of the invention. Although novel, many of the
CC polypeptides of the invention have homology to known proteins, thereby
CC giving an insight into their probable biological activities, and hence
CC potential therapeutic applications. The polypeptides of the invention may
CC have various activities, including cytokine, cell proliferation or cell
CC differentiation activities; stem cell growth factor activity;
CC haematopoiesis regulatory activity; tissue growth activity;
CC immunomodulatory activity; activin- or inhibin-related activities;
CC chemotactic or chemokinetic activities; haemostatic, thrombotic or
CC thrombolytic activities; receptor or ligand activities; or may be
CC involved in oncogenesis, cancer cell proliferation or metastasis.
CC Depending on their biological activities, polypeptides and nucleotides of
CC the invention are useful for preventing, treating or ameliorating medical
CC conditions, e.g., by protein or gene therapy. Such conditions include
CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell
CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),
CC proliferative retinopathy, atherosclerosis, coronary heart disease,
CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal
CC vascular growth. Polypeptides involved with tissue regeneration and
CC repair (or nucleic acids encoding them) may be used to promote wound
CC healing (e.g., of burns, incisions and ulcers), while those with
CC immunomodulatory activities may be used in the treatment of viral,
CC bacterial and fungal infections in addition to immune disorders.
CC Polypeptides with growth factor activity may be used in cell cultures to
CC promote cell growth. For example, such polypeptides may be used to
CC manipulate stem cells in culture to give rise to neuroepithelial cells
CC that can be used to augment or replace cells damaged by illness,
CC autoimmune disease or accidental damage. The polypeptides and nucleotides
CC may also be used in the diagnosis of the above conditions, and in drug
CC screening techniques. The present sequence represents a novel human
CC polypeptide of the invention.

XX Sequence 280 AA;
SQ Query Match 100.0%; Score 1002; DB 22; Length 280;
Best Local Similarity 100.0%; Pred. No. 3.6e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MMNFQPPSKAWRASQMTFFIFLLFPSSFTGVLTATITWRLKPSADCGPFRGLPLFIH 60
Db 91 MMNFQPPSKAWRASQMTFFIFLLFPSSFTGVLTATITWRLKPSADCGPFRGLPLFIH 150
QY 61 SIYSWIDTLSTRPGYLWVWYIYRNLIQSVHFFILTLVLIITYLYWQITGGRKIMIRLL 120
Db 151 SIYSWIDTLSTRPGYLWVWYIYRNLIQSVHFFILTLVLIITYLYWQITGGRKIMIRLL 210
QY 121 HEQIINEGKDKMFLIEKLIKLODMERKANPSSLVLERREVEQOGFLHLGHDGSLDLRSR 180
Db 211 HEQIINEGKDKMFLIEKLIKLODMERKANPSSLVLERREVEQOGFLHLGHDGSLDLRSR 270
QY 181 RSVQEGNPRA 190
Db 271 RSVQEGNPRA 280

RESULT 5

AAM80088
ID AAM80088 standard; Protein; 280 AA.

XX AAM80088;

DT 06-NOV-2001 (first entry)

DE Human protein SEQ ID NO 3734.

KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation.

OS Homo sapiens.

PN WO200157190-A2.

PD 09-AUG-2001.

PF 05-FEB-2001; 2001WO-US04098.

PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.
PR 19-JUL-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.

PA (HYSE-) HYSEQ INC.

PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;

DR WPI; 2001-476283/51.
DR N-PSDB; AAK53221.

PT Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -

PS Claim 20; Page 421; 6221pp; English.

CC The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce

CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
CC inflammation.
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666
CC (AAM80020) are omitted as the relevant pages from the sequence listing
CC were missing at the time of publication.

XX SQ Sequence 280 AA;

Query Match 100.0%; Score 1002; DB 22; Length 280;
Best Local Similarity 100.0%; Pred. No. 3.6e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MMNFQPPSKAWRASQMTFFIFLLFPSSFTGVLTATITWRLKPSADCGPFRGLPLFIH 60

Db 91 MMNFQPPSKAWRASQMTFFIFLLFPSSFTGVLTATITWRLKPSADCGPFRGLPLFIH 150

QY 61 SIYSWIDTLSTRPGYLWVWYIYRNLIQSVHFFILTLVLIITYLYWQITGGRKIMIRLL 120

Db 151 SIYSWIDTLSTRPGYLWVWYIYRNLIQSVHFFILTLVLIITYLYWQITGGRKIMIRLL 210

QY 121 HEQIINEGKDKMFLIEKLIKLODMERKANPSSLVLERREVEQOGFLHLGHDGSLDLRSR 180

Db 211 HEQIINEGKDKMFLIEKLIKLODMERKANPSSLVLERREVEQOGFLHLGHDGSLDLRSR 270

QY 181 RSVQEGNPRA 190

Db 271 RSVQEGNPRA 280

RESULT 6

AAB95481
ID AAB95481 standard; Protein; 330 AA.

XX AAB95481;

DT 26-JUN-2001 (first entry)

DE Human protein sequence SEQ ID NO:18002.

KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.

OS Homo sapiens.

PN EP1074617-A2.

PD 07-FEB-2001.

PF 28-JUL-2000; 2000EP-0116126.

PR 29-JUL-1999; 99JP-0248036.

PR 27-AUG-1999; 99JP-0300253.

PR 11-JAN-2000; 2000JP-0118776.

PR 02-MAY-2000; 2000JP-0183767.

PR 09-JUN-2000; 2000JP-0241899.

PA (HELI-) HELIX RES INST.

PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Makamatsu A, Nagai K, Otsuki T;

DR WPI; 2001-318749/34.

PT Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -

PS Claim 8; SEQ ID 18002; 2537pp + CD ROM; English.
XX
CC The present invention describes primer sets for synthesising 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dt primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesising polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.
XX
SQ Sequence 330 AA;

Query Match 100.0%; Score 1002; DB 22; Length 330;
Best Local Similarity 100.0%; Pred. No. 4.5e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MMNFPPSKAMRASQMTFFIFLLFPSPFTGVLCTLAITIWRLKPSADCGPFRGLPLFIH 60
Db 141 MMNFPPSKAMRASQMTFFIFLLFPSPFTGVLCTLAITIWRLKPSADCGPFRGLPLFIH 200

OY 61 SIYSWIDTLSTRPGYLWVWYIYRNLIIGSVHFFILTLIVLITFYLYWQITTEGRKIMIRLL 120
Db 201 SIYSWIDTLSTRPGYLWVWYIYRNLIIGSVHFFILTLIVLITFYLYWQITTEGRKIMIRLL 260

OY 121 HEQITNEGKDKMFLIEKLKIQDMKKANPSSLVLERREVEQOGFLHGEHDSLDLSR 180
Db 261 HEQITNEGKDKMFLIEKLKIQDMKKANPSSLVLERREVEQOGFLHGEHDSLDLSR 320

OY 181 RSVQEGNPRA 190
Db 321 RSVQEGNPRA 330

RESULT 7
AAB08764
ID AAB08764 standard; Protein; 387 AA.
XX
AC AAB08764;
XX
DT 02-JAN-2001 (first entry)
XX
DE A human leukocyte and blood related protein (LBAP).
XX
XX Human; leukocyte and blood related protein; LBAP; arteriosclerosis;
KW cell proliferative disorder; actinic keratosis; atherosclerosis;
KW bursitis; cirrhosis; hepatitis; mixed connective tissue disease; MCTD;
KW myelofibrosis; paroxysmal nocturnal hemoglobinuria; cancer;
KW adenocarcinoma; leukemia; lymphoma; melanoma; myeloma; sarcoma;
KW teratocarcinoma; autoimmune disorder; inflammatory disorder;
KW acquired immunodeficiency syndrome; AIDS; Addison's disease;
KW adult respiratory distress syndrome; allergy; ankylosing spondylitis;
KW amyloidosis; anaemia; asthma; autoimmune haemolytic anaemia; infection;
KW Werner syndrome; haemodialysis; extracorporeal circulation; trauma.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Peptide 1..51

FT Domain /note= "signal peptide"
FT 74..94
FT /note= "transmembrane domain"
FT Modified-site 101
FT /note= "potential phosphorylation site"
FT Domain 114..134
FT /note= "transmembrane domain"
FT Modified-site 163
FT /note= "potential phosphorylation site"
FT Domain 167..189
FT /note= "transmembrane domain"
FT Modified-site 194
FT /note= "potential glycosylation site"
FT Domain 213..237
FT /note= "transmembrane domain"
FT Modified-site 261
FT /note= "potential phosphorylation site"
FT Modified-site 267
FT /note= "potential phosphorylation site"
FT Domain 281..299
FT /note= "transmembrane domain"
FT Modified-site 376
FT /note= "potential phosphorylation site"
FT Modified-site 379
FT /note= "potential phosphorylation site"

PN WO200052161-A2.
XX
XX 08-SEP-2000.
PD
XX 29-FEB-2000; 2000WO-US05153.
PF
XX 01-MAR-1999; 99US-0122080.
PR
XX (INCY-) INCYTE PHARM INC.
PA
XX Lal P, Yue H, Hillman JL, Lu DAM, Baughn MR, Tang YT, Azimzal Y;
PI
XX WPI: 2000-587310/55.
DR N-PSDB; AAA64684.
DR
XX
PT Leukocyte and blood associated proteins and polynucleotides encoding
PT them, useful for diagnosis, treatment and prevention of
PT autoimmune/inflammatory disorders and cell proliferative disorders
PT including cancer -
XX
XX
PS Claim 1; Page 65; 70pp; English.
XX
XX The present sequence presents a human leukocyte and blood related
CC protein, designated LBAP. LBAP polynucleotides and polypeptides are
CC useful for treating or preventing a disorder associated with decreased
CC expression or activity of LBAP including a cell proliferative disorder
CC such as actinic keratosis, arteriosclerosis, atherosclerosis, bursitis,
CC cirrhosis, hepatitis, mixed connective tissue disease (MCTD),
CC myelofibrosis, paroxysmal nocturnal hemoglobinuria, etc., cancers
CC including adenocarcinoma, leukemia, lymphoma, melanoma, myeloma,
CC sarcoma, teratocarcinoma and in particular cancers of the adrenal
CC gland, bladder, bone, bone marrow, brain, breast, cervix, etc., and
CC an autoimmune/inflammatory disorder such as acquired immunodeficiency
CC syndrome (AIDS), Addison's disease, adult respiratory distress syndrome,
CC allergies, ankylosing spondylitis, amyloidosis, anaemia, asthma,
CC atherosclerosis, autoimmune haemolytic anaemia, etc., Werner syndrome,
CC complications of cancer, haemodialysis, and extracorporeal circulation,
CC viral, bacterial, fungal, parasitic, protozoan, and helminthic
CC infections, and trauma.
XX
SQ Sequence 387 AA;

Query Match 100.0%; Score 1002; DB 21; Length 387;
Best Local Similarity 100.0%; Pred. No. 5.6e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MMNFPPSKAMRASQMTFFIFLLFPSPFTGVLCTLAITIWRLKPSADCGPFRGLPLFIH 60

Db 198 MMNFQPPSKAWRASQMTFFIFLLFPSPFTGVLTATITWRLKPSADCGPFRGLPLFIH 257
QY 61 SIYSWIDTLSTRPGYLWVWYIRNLIGSVHFFELTLIVLIITYLYWQITEGRKIMIRLL 120
Db 258 SIYSWIDTLSTRPGYLWVWYIRNLIGSVHFFELTLIVLIITYLYWQITEGRKIMIRLL 317
QY 121 HEQIINEGDKMFLEIKLIKQDMMEKKANPSSLVLERREVEQOGFLHLGHDGSLDIRSR 180
Db 318 HEQIINEGDKMFLEIKLIKQDMMEKKANPSSLVLERREVEQOGFLHLGHDGSLDIRSR 377
OY 181 RSVQEGNPRA 190
Db 378 RSVQEGNPRA 387

RESULT 8
AAB83081
ID AAB83081 standard; Protein; 438 AA.
XX
AC AAB83081;
XX
DT 29-JUN-2001 (first entry)
XX
DE Human CASB6411-related partial polypeptide #1.
XX
KW Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;
KM ovarian cancer; colon cancer; autoimmune disease.
XX
OS Homo sapiens.
XX
PN WO200123417-A2.
XX
PD 05-APR-2001.
XX
PF 27-SEP-2000; 2000WO-EP09500.
XX
PR 30-SEP-1999; 99GB-0023154.
PR 07-JUL-2000; 2000GB-0016839.
XX
PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
XX
PI Vinals De Bassols YC;
XX
DR WPI: 2001-316133/33.
DR N-PSDB; AAF82462.
XX
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for
PT prophylactic and therapeutic treatment of cancers, particularly ovarian
PT and colon cancers, autoimmune diseases and related conditions -
XX
XX
PS Disclosure; Page 66; 95pp; English.
XX
CC The present sequence is provided in a specification relating
CC to CASB6411 polypeptides comprising a sequence having at least 70%
CC identity to a sequence of 460 or 154 amino acids fully defined in
CC the specification. CASB6411 polypeptides and polynucleotides are
CC useful for treating a subject by immunoprophylaxis or therapy.
CC The CASB6411 polypeptides are useful in diagnostics, and as
CC vaccines for prophylactic and therapeutic treatment of cancers,
CC particularly ovarian and colon cancers, autoimmune diseases and related
CC conditions. CASB6411 polypeptides are also useful for the
CC structure-based design of agonists, antagonists or inhibitors of the
CC polypeptide.
XX
SQ Sequence 438 AA;

Query Match 100.0%; Score 1002; DB 22; Length 438;
Best Local Similarity 100.0%; Pred. No. 6.6e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
*QY 1 MMNFQPPSKAWRASQMTFFIFLLFPSPFTGVLTATITWRLKPSADCGPFRGLPLFIH 60
|||||

Db 249 MMNFQPPSKAWRASQMTFFIFLLFPSPFTGVLTATITWRLKPSADCGPFRGLPLFIH 308
QY 61 SIYSWIDTLSTRPGYLWVWYIRNLIGSVHFFELTLIVLIITYLYWQITEGRKIMIRLL 120
Db 309 SIYSWIDTLSTRPGYLWVWYIRNLIGSVHFFELTLIVLIITYLYWQITEGRKIMIRLL 368
QY 121 HEQIINEGDKMFLEIKLIKQDMMEKKANPSSLVLERREVEQOGFLHLGHDGSLDIRSR 180
Db 369 HEQIINEGDKMFLEIKLIKQDMMEKKANPSSLVLERREVEQOGFLHLGHDGSLDIRSR 428
QY 181 RSVQEGNPRA 190
Db 429 RSVQEGNPRA 438

RESULT 9
AAB83079
ID AAB83079 standard; Protein; 460 AA.
XX
AC AAB83079;
XX
DT 29-JUN-2001 (first entry)
XX
DE Human CASB6411 protein.
XX
KW Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;
KM ovarian cancer; colon cancer; autoimmune disease.
XX
OS Homo sapiens.
XX
PN WO200123417-A2.
XX
PD 05-APR-2001.
XX
PF 27-SEP-2000; 2000WO-EP09500.
XX
PR 30-SEP-1999; 99GB-0023154.
PR 07-JUL-2000; 2000GB-0016839.
XX
PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
XX
PI Vinals De Bassols YC;
XX
DR WPI: 2001-316133/33.
DR N-PSDB; AAF82460.
XX
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for
PT prophylactic and therapeutic treatment of cancers, particularly ovarian
PT and colon cancers, autoimmune diseases and related conditions -
XX
XX
PS Claim 1; Page 64; 95pp; English.
XX
CC The present sequence is human CASB6411 polypeptide. The
CC invention relates to CASB6411 polypeptides comprising a sequence
CC having at least 70% identity to a sequence of 460 or 154 amino acids
CC fully defined in the specification. CASB6411 polypeptides and
CC polynucleotides are useful for treating a subject by immunoprophylaxis
CC or therapy. The CASB6411 polypeptides are useful in diagnostics, and
CC as vaccines for prophylactic and therapeutic treatment of cancers,
CC particularly ovarian and colon cancers, autoimmune diseases and related
CC conditions. CASB6411 polypeptides are also useful for the
CC structure-based design of agonists, antagonists or inhibitors of the
CC polypeptide. The full length mRNA encoding the present sequence may
CC be alternatively spliced to generate a mRNA encoding a truncated
CC CASB6411 protein.
XX
SQ Sequence 460 AA;

Query Match 100.0%; Score 1002; DB 22; Length 460;
Best Local Similarity 100.0%; Pred. No. 7e-111;
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MMNFQPPSKAWRASQMTFFIFLLFPSPFTGVLTATITWRLKPSADCGPFRGLPLFIH 60
|||||

|||||
Db 271 MMNQPSPKAWRASQMMTFITLLFPSPFTGVLCTLAITITWRLKPSADCGPFRGLPLFIH 330
Qy 61 SIYSWIDTLSTRPGYLWVWYIRNLIGSVHFFITLLIYLYTWQITEGRKIMIRLL 120
Db 331 SIYSWIDTLSTRPGYLWVWYIRNLIGSVHFFITLLIYLYTWQITEGRKIMIRLL 390
Qy 121 HEQIINEGDKMFLEIKLQDMKKNPSSLYERREVEOQGFHLGHDGSLDRSR 180
Db 391 HEQIINEGDKMFLEIKLQDMKKNPSSLYERREVEOQGFHLGHDGSLDRSR 450
Qy 181 RSVQEGNPRA 190
Db 451 RSVQEGNPRA 460
RESULT 10
ABP41828
ID ABP41828 standard; Protein: 305 AA.
XX
AC ABP41828;
XX
DT 22-AUG-2002 (first entry)
XX
DE Human ovarian antigen HACMU05, SEQ ID NO:2960.
XX
KW Human; ovarian antigen; ovary; ovarian; breast; cancer; tumour;
KW ovarian cancer; breast cancer; tumour; reproductive system disorder;
KW infertility; pregnancy disorder; anovulation; polycystic ovary syndrome;
KW PCOS; ovarian cyst; dysmenorrhoea; endocrine disorder; infection;
KW inflammatory condition; immune disorder; blood disorder;
KW cardiovascular disorder; respiratory disorder; neurological disorder;
KW gastrointestinal disorder; urinary system disorder; drug screening;
KW gene therapy; chromosome mapping; forensic analysis;
KW antibody preparation; cytostatic; immunomodulatory; neuroprotective;
KW antiinflammatory; gynaecological; reproductive; chromosome 17q25.
XX
OS Homo sapiens.
XX
PN WO200200677-A1.
XX
PD 03-JAN-2002.
XX
PE 07-JUN-2001; 2001WO-US18569.
XX
PR 07-JUN-2000; 2000US-209467P.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Birse CE, Rosen CA;
XX
DR WPI; 2002-147878/19.
DR N-PSDB; ABQ54905.
XX
PT Isolated nucleic acid molecules encoding novel ovarian polypeptides,
PT useful in the prevention, treatment and diagnosis of cancer (e.g.
PT ovarian cancer), immune disorders, cardiovascular disorders and
PT neurological diseases -
XX
PS Claim 11, SEQ ID NO 2960; 2922pp; English.
XX
CC The invention relates to 2175 novel human ovarian antigens (ABP41054-
CC ABP43228) and to cDNAs encoding them (ABQ54131-ABQ56305), and also
CC encompasses polypeptides 90% identical and polynucleotides 95% identical
CC to the sequences of the invention. The invention additionally relates to
CC recombinant vectors and host cells comprising human ovarian antigen
CC polynucleotides, antibodies against human ovarian antigens, and the use
CC of ovarian antigen polynucleotides and polypeptides in diagnosing,
CC treating, prognosing or preventing various ovary and/or breast-related
CC disorders. Such conditions include ovarian cancer and breast cancer, and
CC metastatic tumours of ovarian or breast origin, reproductive system
CC disorders (e.g., infertility, disorders of pregnancy, anovulation,
CC polycystic ovary syndrome, ovarian cysts, and dysmenorrhoea), endocrine

CC disorders, infections (e.g., chlamydia, HIV, toxoplasmosis, and toxic
CC shock syndrome), inflammatory conditions (e.g., mastitis, oophoritis and
CC vaginitis), immune disorders (e.g., congenital and acquired
CC immunodeficiencies, autoimmune oophoritis, systemic lupus erythematosus),
CC blood-related disorders (e.g., anaemia), cardiovascular disorders,
CC respiratory disorders, neurological disorders, gastrointestinal disorders
CC and urinary system disorders. Ovarian antigen polypeptides and
CC polynucleotides may also be used in screening for compounds which
CC modulate ovarian antigen expression or activity. The polynucleotides may
CC further be used for gene therapy, chromosome mapping, in the
CC identification of individuals and in forensic analysis, and the
CC polypeptides may be used as food additives or to prepare antibodies
CC useful in disease diagnosis, drug targeting and phenotyping. The present
CC sequence represents a human ovarian antigen of the invention.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 305 AA;
XX
Query Match 29.2%; Score 292.5; DB 23; Length 305;
Best Local Similarity 34.9%; Pred. No. 2.8e-26;
Matches 60; Conservative 36; Mismatches 63; Indels 13; Gaps 2;
Qy 1 MMNQPSPKAWRASQMMTFITLLFPSPFTGVLCTLAITITWRLKPSADCGPFRGLPLFIH 60
Db 133 LANQAPRRPWLASHMSTVFITLLCFPAFLGAAYFLCYAWQVKPSSTCGPFRTDITMYE 192
Qy 61 SIYSWIDTL-STRPGYLWVWYIRNLIGSVHFFITLLIYLYTWQITEGRKIMIRL 119
Db 193 AGRWVVRHLAAGPRYSWLPWVHRXYMENTFFVFLVSALLAVIYLNQVVRGQKVICL 252
Qy 120 LHEQIINEGDKMFLEIKLQD-----MEKKNPSSLYERRE 159
Db 253 LKEQISNEGDKIFLINKLSHYERKERERSRVGTTEAAAPALITDEOD 304
RESULT 11
ABB39891
ID ABB39891 standard; Peptide: 31 AA.
XX
AC ABB39891;
XX
DT 04-FEB-2002 (first entry)
XX
DE Peptide #7397 encoded by human foetal liver single exon probe.
XX
KW Human; foetal liver; gene expression; single exon nucleic acid probe.
XX
OS Homo sapiens.
XX
PN WO200157277-A2.
XX
PD 09-AUG-2001.
XX
PE 30-JAN-2001; 2001WO-US00669.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MODE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI; 2001-483447/52.
XX
PT Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human fetal liver -

XX
PS Claim 27; SEQ ID NO 32526; 639pp + sequence listing; English.
XX
CC The invention relates to a single exon nucleic acid probe for
CC measuring human gene expression in a sample derived from human foetal
CC liver. The single exon nucleic acid probes may be used for predicting,
CC measuring and displaying gene expression in samples derived from human
CC fetal liver. The present sequence is a peptide encoded by a single exon
CC nucleic acid probe of the invention.
CC Note: The sequence data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 31 AA;

Query Match 14.8%; Score 148; DB 22; Length 31;
Best Local Similarity 100.0%; Pred. No. 2.6e-10;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 131 KMFLIEKLIKLDMEKKANPSSLYERREVE 161
Db 1 KMFLIEKLIKLDMEKKANPSSLYERREVE 31

RESULT 12
AAM60631
ID AAM60631 standard; Protein; 31 AA.
XX
AC AAM60631;
XX
DT 05-NOV-2001 (first entry)
XX
DE Human brain expressed single exon probe encoded protein SEQ ID NO: 32736.
XX
KW Human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer.
XX
OS Homo sapiens.
XX
PN WO200157275-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00667.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI; 2001-483446/52.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
PT brains -
XX
PS Example 4; SEQ ID NO: 32736; 650pp + Sequence Listing; English.
XX
CC The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is a protein encoded by one of
CC the probes of the invention.
XX

SQ Sequence 31 AA;

Query Match 14.8%; Score 148; DB 22; Length 31;
Best Local Similarity 100.0%; Pred. No. 2.6e-10;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 131 KMFLIEKLIKLDMEKKANPSSLYERREVE 161
Db 1 KMFLIEKLIKLDMEKKANPSSLYERREVE 31

RESULT 13
AAM73303
ID AAM73303 standard; Protein; 31 AA.
XX
AC AAM73303;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human bone marrow expressed probe encoded protein SEQ ID NO: 33609.
XX
KW Human; bone marrow expressed exon; gene expression analysis; probe;
KW microarray; cancer; leukaemia; lymphoma; myeloma.
XX
OS Homo sapiens.
XX
PN WO200157276-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00668.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI; 2001-488900/53.
XX
PT Human genome-derived single exon nucleic acid probes useful for
PT analyzing gene expression in human bone marrow -
XX
PS Example 4; SEQ ID NO: 33609; 658pp + Sequence Listing; English.
XX
CC The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC bone marrow. They can be used to measure gene expression in bone marrow
CC samples, which may enable the improved diagnosis and treatment of cancers
CC such as lymphoma, leukaemia and myeloma. The present sequence is a
CC protein encoded by one of the probes of the invention.
XX
SQ Sequence 31 AA;

Query Match 14.8%; Score 148; DB 22; Length 31;
Best Local Similarity 100.0%; Pred. No. 2.6e-10;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 131 KMFLIEKLIKLDMEKKANPSSLYERREVE 161
Db 1 KMFLIEKLIKLDMEKKANPSSLYERREVE 31

RESULT 14
AAM33503
ID AAM33503 standard; Protein; 31 AA.
XX

```

AC AAM33503;
XX
XX 17-OCT-2001 (first entry)
XX
XX Peptide #7540 encoded by probe for measuring placental gene expression.
DE
XX Probe; microarray; human; placenta; antenatal diagnosis;
XX genetic disorder.
XX
XX Homo sapiens.
XX
XX WO200157272-A2.
XX
XX 09-AUG-2001.
XX
XX 30-JAN-2001; 2001WO-US00663.
XX
XX 04-FEB-2000; 2000US-0180312.
XX
XX 26-MAY-2000; 2000US-0207456.
XX
XX 30-JUN-2000; 2000US-0608408.
XX
XX 03-AUG-2000; 2000US-0632366.
XX
XX 21-SEP-2000; 2000US-0234687.
XX
XX 27-SEP-2000; 2000US-0236359.
XX
XX 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI; 2001-488897/53.
XX
XX Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human placenta -
XX
XX Claim 27; SEQ ID No 33772; 654bp; English.
XX
XX The present invention relates to single exon nucleic acid probes (SENP:
XX see AAI3315-AA157546). The present sequence is a peptide encoded by one
XX such probe. The probes are useful for producing a microarray for
XX predicting, measuring and displaying gene expression in samples derived
XX from human placenta. The probes are useful for antenatal diagnosis of
XX human genetic disorders.
XX
XX Sequence 31 AA;
SQ
Query Match 14.8%; Score 148; DB 22; Length 31;
Best Local Similarity 100.0%; Pred. No. 2.6e-10;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 131 KMFLIEKLKLODMKKANPSSLYLERREVE 161
DB 1 KMFLIEKLKLODMKKANPSSLYLERREVE 31
RESULT 15
ABG43154
ID ABG43154 standard; Peptide; 31 AA.
AC ABG43154;
XX
XX 19-AUG-2002 (first entry)
XX
XX Human peptide encoded by genome-derived single exon probe SEQ ID 32819.
XX
XX Human; single exon probe; asthma; lung cancer; COPD; ILD;
XX chronic obstructive pulmonary disease; interstitial lung disease;
XX familial idiopathic pulmonary fibrosis; neurofibromatosis;
XX tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
XX Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
XX pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndrome;
XX pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
XX primary ciliary dyskinesia; pulmonary hypertension;
XX hyaline membrane disease.

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XX
XX Homo sapiens.
XX
XX WO200186003-A2.
XX
XX 15-NOV-2001.
XX
XX 30-JAN-2001; 2001WO-US00665.
XX
XX 04-FEB-2000; 2000US-180312P.
XX
XX 26-MAY-2000; 2000US-207456P.
XX
XX 30-JUN-2000; 2000US-0608408.
XX
XX 03-AUG-2000; 2000US-0632366.
XX
XX 21-SEP-2000; 2000US-234687P.
XX
XX 27-SEP-2000; 2000US-236359P.
XX
XX 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX WPI; 2002-114183/15.
XX
XX Spatially-addressable set of single exon nucleic acid probes, used to
XX measure gene expression in human lung samples -
XX
XX Claim 27; SEQ ID No 32819; 634bp; English.
XX
XX The invention relates to a spatially-addressable set of single exon
XX nucleic acid probes for measuring gene expression in a sample derived
XX from human lung comprising single exon nucleic acid probes having one of
XX 12614 nucleic acid sequences mentioned in the specification, or their
XX complements or the 12387 open reading frames derived from the 12614
XX probes. Also included are a microarray comprising the novel set of
XX probes; the novel set of probes which hybridise at high stringency to a
XX nucleic acid expressed in the human lung; measuring gene expression in a
XX sample derived from human lung, comprising (a) contacting the array with
XX a collection of detectably labeled nucleic acids derived from human lung
XX mRNA, and (b) measuring the label detectably bound to each probe of
XX the array; identifying exons in a eukaryotic genome, comprising
XX (a) algorithmically predicting at least one exon from genomic sequences
XX of the eukaryote; and (b) detecting specific hybridisation of detectably
XX labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
XX having a fragment identical to the predicted exon, the probe is included
XX in the above mentioned microarray; assigning exons to a single gene,
XX comprising (a) identifying exons from genomic sequence by the method
XX above and (b) measuring the expression of each of the exons in several
XX tissues and/or cell types using hybridisation to a single exon
XX microarrays having a probe with the exon, where a common pattern of
XX expression of the exons in the tissues and/or cell types indicates that
XX the exons should be assigned to a single gene; a peptide comprising one
XX of 12011 sequences, mentioned in the specification, or encoded by the
XX probes/open reading frames (ORF). The probes are used for gene
XX expression analysis, and for identifying exons in a gene, particularly
XX using human lung derived mRNA and for the study of lung diseases
XX such as asthma, lung cancer, chronic obstructive pulmonary disease
XX (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
XX fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
XX Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
XX haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
XX pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic
XX pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
XX and hyaline membrane disease. The present sequence is a peptide/protein
XX encoded by a single exon probe of the invention.
XX Note: The sequence data for this patent did not form part
XX of the printed specification, but was obtained in electronic
XX format directly from WPI at
XX ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 31 AA;
SQ
Query Match 14.8%; Score 148; DB 23; Length 31;
Best Local Similarity 100.0%; Pred. No. 2.6e-10;

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Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 131 KMFLIEKLIKLODMCKKANPSSLYLRRREVE 161
|||||
Db 1 KMFLIEKLIKLODMCKKANPSSLYLRRREVE 31

Search completed: November 9, 2002, 04:30:32
Job time : 81 secs